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(21) International Application Number: PCT/CA98/01069 (22) International Filing Date: 18 November 1998 (18.11.98) (30) Priority Data: 60/065,692 18 November 1997 (18.11.97) US (71)(72) Applicants and Inventors: MOSKOVITS, Martin [CA/CA]; 145 Chiltern Hill Road, Toronto, Ontario M6C 3C3 (CA). LI, Jing [CN/CA]; Apartment 1504, 255 Glenlake Avenue, Toronto, Ontario M6P 1G2 (CA). HASLETT, Thomas, L. [CA/CA]; 9 Redwood Avenue, Toronto, Ontario M4L 2S5 (CA). (74) Agents: SCHUMACHER, Lynn, C. et al. ; Hill & Schumacher, Suite 802, 335 Bay Street, Toronto, Ontario M5H 2R3 (CA).	(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>	
(54) Title: CONTROLLED SYNTHESIS AND METAL-FILLING OF ALIGNED CARBON NANOTUBES (57) Abstract <p>The present invention provides a powerful new method for producing, uniform sized and uniformly aligned nanotubes through catalytic pyrolysis of a hydrocarbon within the dense, uniform and parallel pores of alumina nano-templates. The catalyst, Co, Fe, Ni or another suitable substance is deposited electrochemically into the bottom of the channel of the alumina template. The nanotubes with any desired diameter in the range 5-500 nm and lengths up to ~ 100 µm, are generated by the pyrolysis of a suitable hydrocarbon inside the pores of the alumina template with at least one end open at the alumina/air interface. The nanotubes may be filled by metals using for example electroless deposition.</p>		

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CONTROLLED SYNTHESIS AND METAL-FILLING OF ALIGNED CARBON NANOTUBES

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FIELD OF THE INVENTION

The present invention relates to a method of synthesis of aligned carbon nanotubes through catalytic pyrolysis of hydrocarbon within alumina nano-templates. The present invention also relates to a method of filling the inner cavity of the nanotubes with metals and more particularly the method relates to electroless plating of metals including but not limited to Ni, Co, their alloys and other metals into the inner cavity of the nanotubes and filling the metals inside.

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BACKGROUND OF THE INVENTION

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Carbon nanotube, a novel carbon material, was theoretically predicted and experimentally shown to possess unique electronic and mechanical properties and is of great scientific and potential commercial value. Recent studies reveal that the diameter, number of concentric cylinders and helicity of the arrangement of the carbon atoms in the nanotube shell determine whether the nanotube has metallic, semiconducting or insulating properties. Hence a wide range of materials properties may be "tuned in" by changing the structural properties of these nanotubes. The stiffness and high strength of these carbon nanotubes make them ideal probes in scanning probe microscopy, and efficient field-emitters, as recently shown by Smalley's group. Additionally the inner hollow cavity of the nanotubes may serve as nanometer scale test tubes in which to carry out interesting experiments with submicroscopic quantities of reagents. Nanotubes filled with metallic or semiconducting particles might serve as the constituents of novel materials with useful magnetic, electrical or electronic properties leading to new devices.

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The conventional synthesis methods for carbon nanotube include

carbon arc discharge (S. Iijima, Nature, **354**, 56, 1991) and catalytic pyrolysis of hydrocarbon (M. Endo, K. Takeuchi, S. Igarashi, K. Kobori, M. Shiraishi, H. W. Kroto, J. Phys. Chem. Solids, **54**, 1841, 1993), which generate nanotubes often containing traces of the catalyst particles used to generate them and possessing highly variable dimensions. Synthesis of aligned nanotube by pyrolysis of hydrocarbon with a patterned cobalt catalyst on silica substrate was recently reported by M. Terrones et al., Nature, **388**, 52, 1997 and with iron nanoparticles in mesoporous silica by W. Z. Li et al., Science, **274**, 1701, 1996. The successful production of carbon nanotubes in an alumina template by pyrolysis of propylene has been disclosed by T. Kyotani, L. Tsai, A. Tomita, Chem. Mater., **8**, 2190, 1996.

The encapsulation of foreign materials in the carbon nanotubes has been disclosed by P. M. Ajayan and S. Iijima (United State Patent, No 5,457,343). Ni, Fe, Co, rare earth metals and their carbides were found to be encapsulated in the nanotubes during synthesis of them in carbon arc discharge by Y. Saito et al., J. Phys. Chem. Solids, **54**, 1849, 1993, and also by C. Guerret-Piecutot et al., Nature, **372**, 761, 1994. The metals with a low melting point, such as lead, zinc, selenium, and molten metal salt, AgNO₃ were filled through capillarity into the carbon nanotubes, reported by P. M. Ajayan et al., Nature, **362**, 522, 1993, E. Dujardin et al., Science **265**, 1850, 1994 and by D. Ugarte et al., Science, **274**, 1897, 1996.

SUMMARY OF THE INVENTION

It is an object of the present to provide a method of producing carbon nanotubes of uniform size and uniformly aligned in the axial direction.

The present invention provides a powerful new method for producing, uniform sized and uniformly aligned nanotubes through catalytic pyrolysis of a hydrocarbon within the dense, uniform and parallel pores of alumina nano-templates. The catalyst, Co, Fe, Ni or another suitable substance

is deposited electrochemically into the bottom of the channel of the alumina template. The nanotubes with any desired diameter in the range 5-500 nm and lengths up to ~100 μ m, are generated by the pyrolysis of a suitable hydrocarbon inside the pores of the alumina template with at least one end open at the alumina/air interface.

The present invention provides a process for filling the carbon nanotubes obtained by above method with metal Ni, Co, Fe, Ag and their alloys by electroless (chemical) plating.

In one aspect of the invention there is provided a process for synthesis of carbon nanotube. The method comprises anodizing an aluminum substrate in an effective bath to produce an alumina template with a plurality of pores each having a pore diameter. The method includes depositing an effective catalyst into the pores and exposing the alumina template with the catalyst containing pores to an effective hydrocarbon gas at an effective temperature to grow carbon nanotubes in the pores. Each carbon nanotube has an outer diameter corresponding to the pore diameter in the template in which the carbon nanotube is produced.

In another aspect of the invention there is provided a process for producing metal containing carbon nanotubes, comprising:

a process for producing metal containing carbon nanotubes, comprising:
anodizing an aluminum substrate in an effective bath to produce an alumina template with a plurality of pores each having a pore diameter;
depositing an effective catalyst into the bottom of the pores;
exposing the alumina template with the catalyst containing pores to an effective hydrocarbon gas at an effective temperature to grow carbon nanotubes in the pores, each carbon nanotube having an outer diameter corresponding to the pore diameter in the template in which the carbon nanotube is produced; and
depositing a metal into the nanotubes.

In this aspect of the invention the metal may be deposited into the

nanotube by electroless deposition.

BRIEF DESCRIPTION OF THE DRAWINGS

5 The process of growth of carbon nanotubes and metal filling of the nanotubes will now be described, by way of example only, reference being had to the accompanying drawings, in which:

Figure 1 is a schematic diagram showing the steps of synthesising carbon nanotubes in accordance with the present invention;

10 Figure 2 is a scanning electron micrograph (SEM) of exposed carbon nanotubes after etching partially away the template surface in an NaOH solution;

Figure 3a shows an electron diffraction pattern of carbon nanotubes produced according to the present invention;

15 Figure 3b shows an electron diffraction pattern of nickel filled carbon nanotubes;

Figure 4a shows a transmission electron micrograph (TEM) of carbon nanotubes produced by the present method;

Figure 4b shows a TEM of nickel filled carbon nanotubes;

20 Figure 5a shows transmission electron micrographs of Ni-P alloy encapsulated within the carbon nanotubes;

Figures 5b and 5d show high resolution images of the interface of Ni-P alloy and carbon; and

Figure 5c shows a high resolution TEM of Ni-P alloy.

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DETAILED DESCRIPTION OF THE INVENTION

The process of synthesizing carbon nanotubes is showed schematically in Figure 1. Anodic oxidation of an aluminum (99.99%) substrate 10 was carried out in 0.3 M of oxalic acid bath at cell voltages of 25, 40, 50 and 65 V respectively at 18° C. While the oxalic acid bath is preferred, other

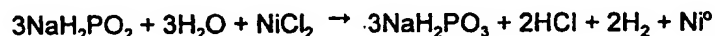
anodizing electrolyte media such as sulphuric, phosphoric and chromic acids may also be used. In these examples the film thickness was controlled to lie in the range 4-10 μm by selecting the anodic oxidation time appropriately. Under the present experimental conditions, the film thickness is calculated using the formula: $D = \alpha t / A$; where α is an experimentally determined constant equal to 0.017 $\mu\text{m}/(\text{mA cm}^2 \text{ min})$; I is the anodizing current in mA; t is the anodizing time in minutes, and A is the aluminum sample area in cm^2 . Films in the range 1-100 μm may be generated in this way. The anodized alumina templates 12 were immersed in 0.1 M phosphoric acid at 30° for 30 min. to widen the pores and to thin the barrier layer. After widening, the average pore diameter of the pores 14 in template 12 obtained at cell voltages of 25, 40, 50 and 65 V are 20, 35, 50 and 70 nm, respectively. The pores 14 are straight, uniform and parallel as shown in the diagrammatic representation in Figure 1.

A cobalt (or iron) catalyst sample 16 approximately 200 nm in length was electrochemically deposited at the bottom of the pores using 14 V and 100 Hz of AC cell voltage for 30 seconds in a bath consisting of 240 g/l of $\text{CoSO}_4 \cdot 7\text{H}_2\text{O}$, (or 120 g/l of $\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$) 40 g/l of HBO_3 and 1 g/l of ascorbic acid. The cobalt-containing templates 12 were placed in a tube furnace and reduced at 600° C for 4-5 hours in flowing CO (100 ml/min). Then a mixture of 10% acetylene in nitrogen was introduced into the reactor at a flow rate of 100 ml/min. Acetylene was decomposed by pyrolysis in the temperature range 640° C to 700° C to form the carbon nanotubes 20 in the template channels. While this is a preferred temperature range the acetylene may be decomposed in a range from 600° C to about 800° C. In addition to acetylene, it will be understood that other hydrocarbon containing gases may be used, for example propylene and methane to mention just a few.

After formation, the nanotubes 20 were annealed to further graphitize the tubes at 700° C in flowing nitrogen for 15 hours. Figure 2 shows SEM image of the nanotubes whose ends were exposed by partially etching the

alumina template with a NaOH solution. The nanotubes 20 are perpendicular to the template 12. Their ends at the alumina/air interface are open and their outer diameters correspond to the inner diameters of the pores 14. An electron diffraction pattern (Figure 3a) recorded for a bundle of the nanotubes that had been released from the alumina template by dissolving the latter in a NaOH solution, (Figure 4a) reveals an ordered graphitic arrangement within the bundles. The inter-layer spacing in the 002 direction (d_{002}) deduced from the electron diffraction data is approximately 3.45 nm, which is close that value in graphite ($d_{002} = 3.35$ nm).

Electroless nickel deposits were produced in an acid bath consisting of 30 g/l of nickel chloride, 10 g/l of sodium hypophosphite, 50 g/l of sodium acetate and ammonia to pH = 5. The alumina templates containing carbon nanotubes were washed with acetone then with distilled water. The washed templates were dipped in the electroless deposition bath for 20-30 min at 25-40° C. The reaction producing the nickel deposit can be formulated as:



It was found that addition of 0.5 g/l of quinhydrone improved the deposition. The quinhydrone may function as a surfactant to increase the wetting of the walls of nanotubes by the solution and as a pH stabilizing agent. Other surfactants, such as gelatine, Dodecyl sulfate sodium salt, are also anticipated to improve the wetting efficacy of the deposition solution. After completely dissolving the alumina template using 0.1 M NaOH solution at 60- 80° C, the metal-containing nanotubes (Figure 4b) could be separated as an insoluble precipitate. An electron diffraction pattern (Figure 3b) recorded for the bundle of nickel-containing nanotubes exhibits clear diffraction spots corresponding to Nickel metal, which coincidentally almost overlaps the diffraction rings produced by the carbon layers of the nanotubes due to the near-correspondence of some of the lattice constants. A lattice constant of approximately 3.52 nm is obtained from

the electron diffraction pattern of the deposited metal. The high-resolution TEM image (Figure 5) shows that the deposited metal is highly crystalline. An EDX analysis of the metal encapsulated in the nanotubes shows it to be a nickel/phosphorous mixture with an average Ni/P ratio of 94% Ni and 6% P.

5 While nickel, cobalt and iron are preferred catalysts for deposition into the pores of the anodized aluminum template for growth of the carbon nanotubes, it is anticipated by the inventors that other suitable or effective catalysts may be used. For example, the metals below each of these three preferred metals are reasonably expected to work to some degree.

10 Similarly, those skilled in the art will appreciate that numerous other metals may be deposited into the carbon nanotubes once formed. Examples include metals such as Ag, Cu, Bi, Pb and Sn may be deposited into the nanotubes by electroless deposition. It will also be understood by those skilled in the art that the nanotubes may be filled with the metals by
15 electrochemical deposition.

 The present method is highly advantageous over prior art methods of growing carbon tubes since in prior art methods the lack of catalyst in the pores yields tubes that exhibit poor uniformity along the tube axis due to the influence of diffusion of the gaseous reagents into and out of the pores of the
20 template.

 In the present method, it is not necessary to strip off the oxide film from the aluminum plate so that a thin film, i.e. a film that is not self-supporting, can also be used as a template allowing one to make tubes of any desired length, by using templates of the appropriate thickness in the practical range 0.1
25 µm to over 100 µm, in contrast to the method described by Kyotani, that requires self-supporting templates and hence is incapable of making tubes below a minimum length below which the template is no longer self-supporting. The present method also provides a much more economical method of growing nanotubes since the number of procedural steps is decreased. The carbon

nanotubes resulting from catalytic, thermal decomposition result in the growth of much more uniform pores in the tubes in contrast to tubes grown in templates without catalyst.

5 The present invention provides a method of producing metal-loaded carbon nanotubes having industrial utility in many areas, including but not limited to catalysis, energy production such as battery technology and may be used as constituents of novel materials with useful electrical, magnetic or electronic properties leading to new devices.

10 The foregoing description of the preferred embodiments of the process for producing carbon nanotubes has been presented to illustrate the principles of the invention and not to limit the invention to the particular embodiments disclosed. It is intended that the scope of the invention be defined by all of the embodiments encompassed within the following claims and their equivalents.

15

WE CLAIM

1. A process for synthesis of carbon nanotubes, comprising:
anodizing an aluminum substrate in an effective bath to produce an alumina template with a plurality of pores each having a pore diameter;
depositing an effective catalyst into the pores; and
exposing said alumina template with the catalyst containing pores to an effective hydrocarbon gas at an effective temperature to grow carbon nanotubes in said pores.
2. The process according to claim 1 wherein the aluminum substrate is anodized under conditions effective to produce said plurality of pores substantially parallel to each other.
3. The process according to claims 1 or 2 wherein said aluminum substrate is anodized in an effective acid bath.
4. The process according to claims 1, 2 or 3 wherein said catalyst is selected from the group consisting of Co, Fe, Ni and alloys thereof.
5. The process according to claims 1, 2, 3 or 4 wherein said catalysts are electrochemically deposited into said pores.
6. The process according to claims 1, 2, 3, 4 or 5 wherein said hydrocarbon gas comprises acetylene.
7. The process according to claim 6 wherein said hydrocarbon gas is pyrolyzed in a temperature range from about 600° C to about 800° C.

8. The process according to claims 1, 2, 3, 4, 5, 6 or 7 wherein subsequent to producing said carbon nanotubes, said carbon nanotubes are annealed to further graphitize said nanotubes.
9. The process according to claims 1, 2, 3, 4, 5, 6, 7 or 8 including depositing a metal into said nanotubes.
10. The process according to claim 9 wherein said metal deposited into said nanotubes is selected from the group consisting of Ni, Fe, Co and Ag.
11. The process according to claim 10 wherein said metal is deposited into said nanotubes by electroless deposition from an effective electroless deposition bath.
12. The process according to claim 10 wherein said metal is deposited into said nanotubes by electrochemical deposition.
13. The process according to claim 11 wherein said electroless deposition bath comprises a surfactant selected from the group consisting of quinuclidine, gelatine and dodecyl sulfate sodium salts as additives for improving the electroless deposition of said metal.
14. The process according to claims 1, 2, 3, 4, 5, 6, or 7 including immersing the alumina substrate in an effective solution to widen said pores and thin an alumina barrier layer on the interior surface of said pores
15. The process according to claim 11 wherein said nanotubes are washed in an effective solution prior to electroless deposition of said metals.

16. The process according to claims 1, 2, 3, 4, 5, 6, 7, 8 or 9 including dissolving the alumina template to separate the carbon nanotubes from said template.
17. The process according to claim 16 wherein said alumina template is dissolved in an NaOH solution.
18. The process according to claim 17 wherein the metal-containing nanotubes are separated as an insoluble precipitate.
19. A process for producing metal containing carbon nanotubes, comprising:
anodizing an aluminum substrate in an effective bath to produce an alumina template with a plurality of pores each having a pore diameter;
depositing an effective catalyst into the bottom of the pores;
exposing said alumina template with the catalyst containing pores to an effective hydrocarbon gas at an effective temperature to grow carbon nanotubes in said pores, each carbon nanotube having an outer diameter corresponding to the pore diameter in the template in which said carbon nanotube is produced;
and
depositing a metal into said nanotubes.
20. The process according to claim 19 wherein said metal is deposited into said nanotube by electroless deposition.
21. The process according to claims 19 or 20 wherein said catalyst is selected from the group consisting of Ni, Co and Fe.
22. The process according to claims 19, 20 or 21 wherein said metal deposited into said nanotubes by electroless deposition includes but is

not limited to Ni, Co, Fe, Ag and their alloys.

23. The process according to claims 19, 20, 21 or 22 wherein said metal deposited into said nanotubes by electroless deposition includes Ag, Cu, Bi, Pb and Sn.

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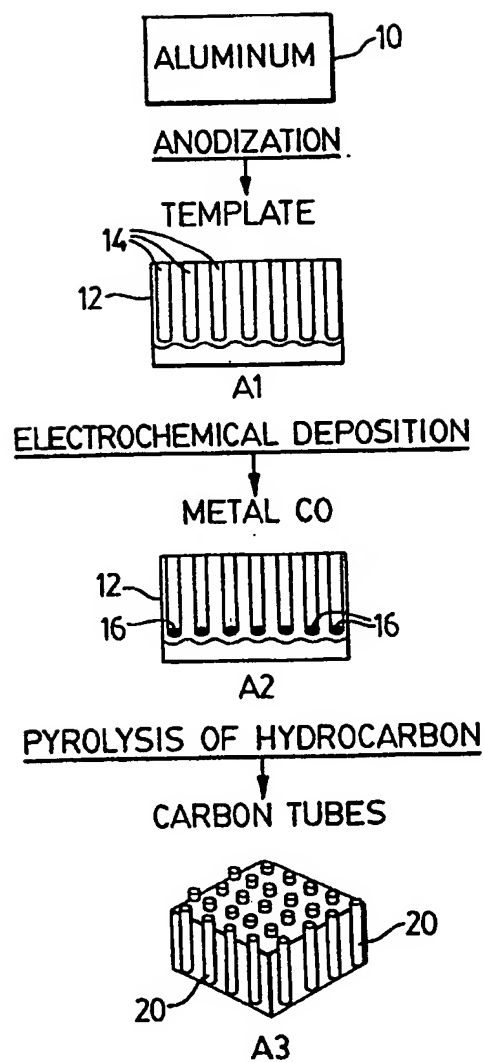


FIG. 1

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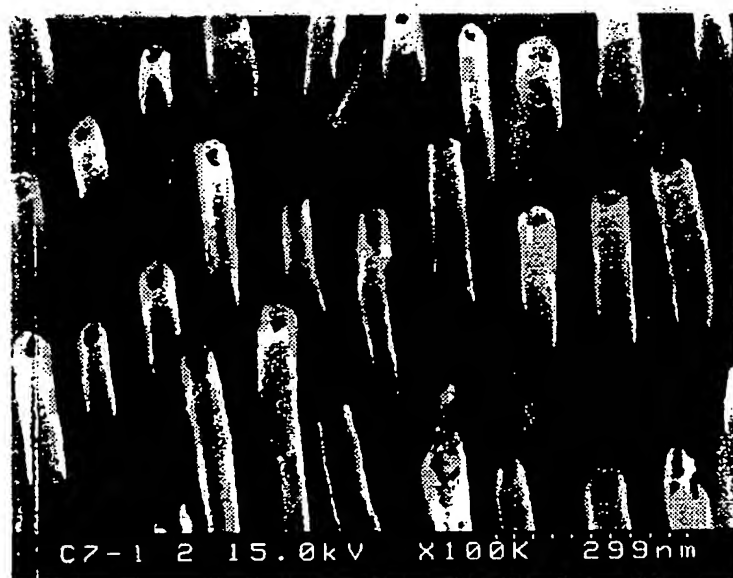


FIG. 2

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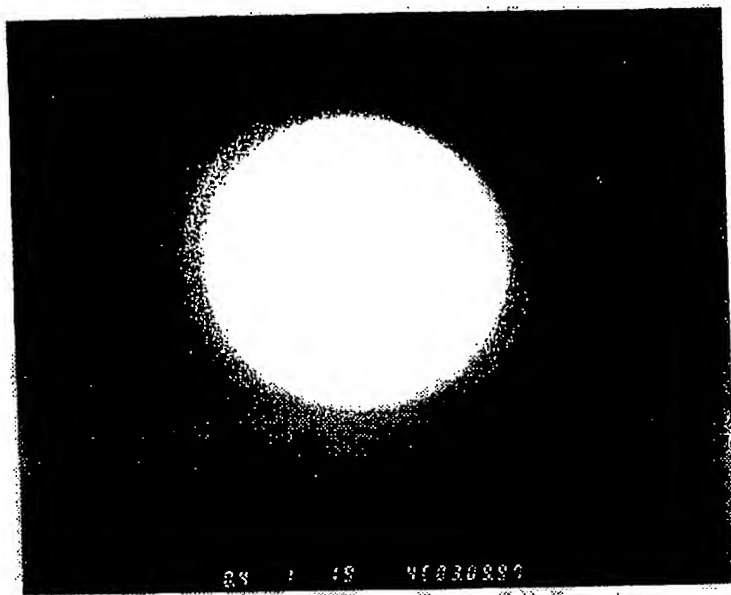


FIG. 3 (a)

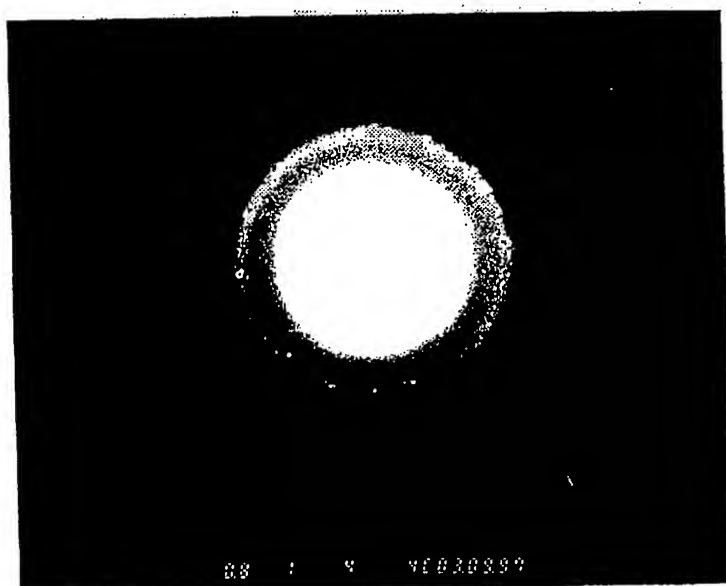


FIG. 3 (b)

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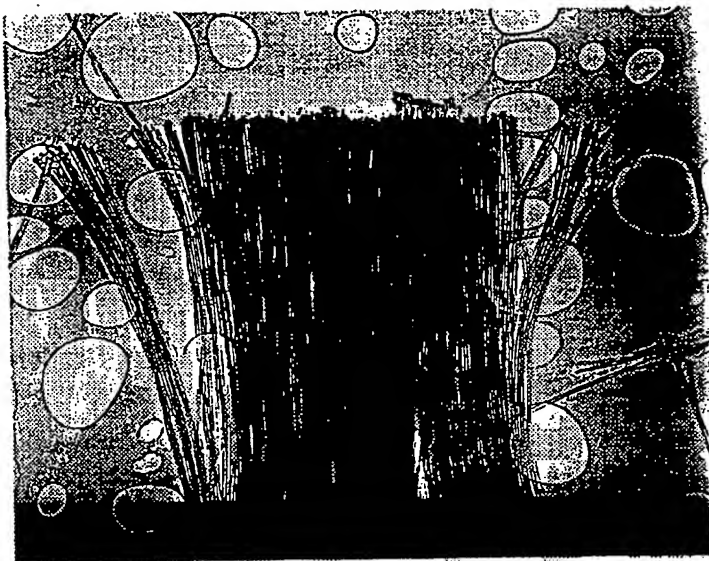


FIG. 4 (a)

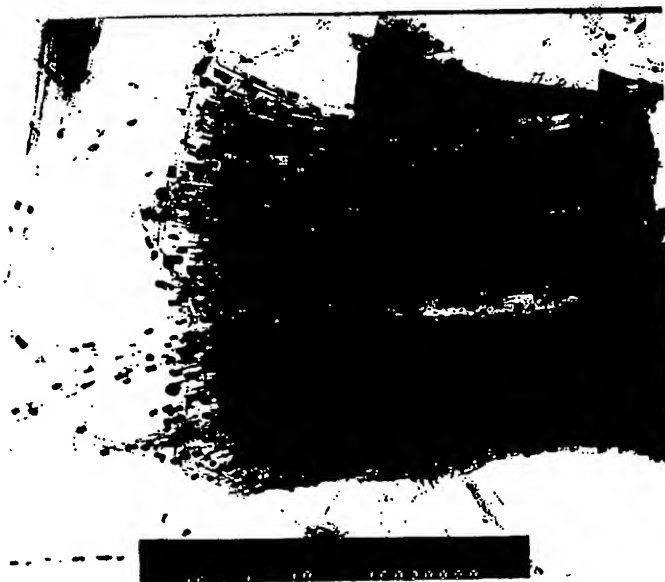
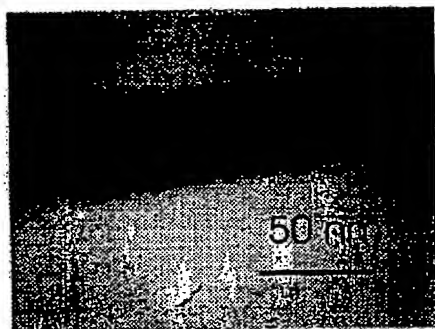


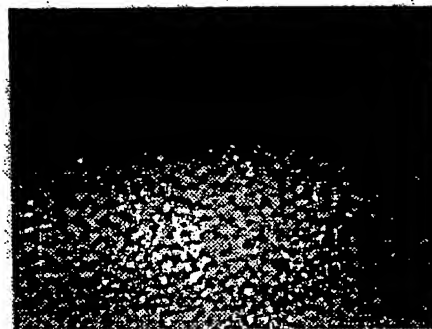
FIG. 4 (b)

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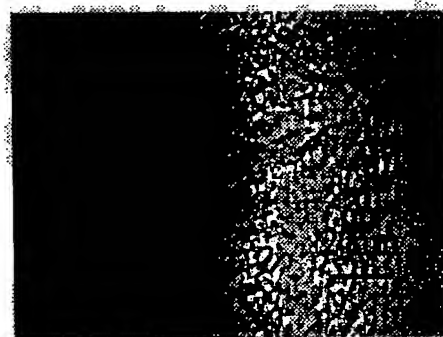
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d

FIG. 5

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INTERNATIONAL SEARCH REPORT

International Application No
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A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 C01831/02

According to International Patent Classification (IPC) or to both national classification and IPC

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Minimum documentation searched (classification system followed by classification symbols)
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Electronic data base consulted during the International search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,X	JING LI ET AL.: "Nanoscale electroless metal deposition in aligned carbon nanotubes" CHEMISTRY OF MATERIALS, vol. 10, no. 7, July 1998, pages 1963-1967, XP002097102 see the whole document	1-11, 13, 15-22
P,X	CHE G ET AL: "CHEMICAL VAPOR DEPOSITION BASED SYNTHESIS OF CARBON NANOTUBES AND NANOFIBERS USING A TEMPLATE METHOD" CHEMISTRY OF MATERIALS, vol. 10, no. 1, January 1998, pages 260-267, XP000776644 see the whole document	1, 4, 8, 16, 18
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INTERNATIONAL SEARCH REPORT

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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	MARTIN CHARLES R: "Nanomaterials: a membrane-based synthetic approach" SCIENCE; SCIENCE DEC 23 1994 AMERICAN ASSOC FOR THE ADVANCEMENT OF SCIENCE, WASHINGTON, DC, USA, vol. 266, no. 5163, 23 December 1994, pages 1961-1966, XP002097060 see page 1964, left-hand column, line 48 - line 52	5
Y	KYOTANI TAKASHI ET AL.: "Formation of platinum nanorods and nanoparticles in uniform carbon nanotubes prepared by a template carbonization method" CHEMICAL COMMUNICATIONS., no. 7, 7 April 1997, pages 701-702, XP002095459 CIETY OF CHEMISTRY GB see the whole document	2,3
A		1,6,7, 9-11,16, 18-20,22
A	KYOTANI TAKASHI ET AL.: "Formation of ultrafine carbon tubes by using an anodic aluminum oxide film as a template" CHEMISTRY OF MATERIALS., vol. 7, no. 8, August 1995, pages 1427-1428, XP002097059 WASHINGTON US see the whole document	1-3,16, 18-20
A	KYOTANI T ET AL: "PREPARATION OF ULTRAFINE CARBON TUBES IN NANOCHANNELS OF AN ANODIC ALUMINUM OXIDE FILM" CHEMISTRY OF MATERIALS, vol. 8, no. 8, August 1996, pages 2109-2113, XP000626894 cited in the application see page 2109, 2110 "Experimental section" see page 2113 "Conclusions"	1-3,16, 18,19
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